



**PRIORITY
DOCUMENT**
SUBMITTED OR TRANSMITTED IN
COMPLIANCE WITH RULE 17.1(a) OR (b)

The Patent Office
Concept House
Cardiff Road
Newport
South Wales
NP10 8QQ

I, the undersigned, being an officer duly authorised in accordance with Section 74(1) and (4) of the Deregulation & Contracting Out Act 1994, to sign and issue certificates on behalf of the Comptroller-General, hereby certify that annexed hereto is a true copy of the documents as originally filed in connection with the patent application identified therein.

In accordance with the Patents (Companies Re-registration) Rules 1982, if a company named in this certificate and any accompanying documents has re-registered under the Companies Act 1980 with the same name as that with which it was registered immediately before re-registration save for the substitution as, or inclusion as, the last part of the name of the words "public limited company" or their equivalents in Welsh, references to the name of the company in this certificate and any accompanying documents shall be treated as references to the name with which it is so re-registered.

In accordance with the rules, the words "public limited company" may be replaced by p.l.c., plc, P.L.C. or PLC.

Re-registration under the Companies Act does not constitute a new legal entity but merely subjects the company to certain additional company law rules.

REC'D 13 JAN 2005	
WIPO	PCT

Signed

AmBraser

Dated 6 January 2005

BEST AVAILABLE COPY

Patents Form 1/77

Patents Act 1977
(Rule 16)

Request for grant of a patent FAX

(See the notes on the back of this form. You can also get an explanatory leaflet, from the Patent Office to help you fill in this form.)

The
Patent
Office

1/77

04 FEB 2004

The Patent Office

Cardiff Road
Newport
South Wales NP10 8QX

1. Your reference

LPT0617

0402471.7

2. Patent application number
(The Patent Office will fill in this part)

3. Full name, address and postcode of the or of each applicant (underline all surnames)

Gursharan Singh Chana
Kenwood
57 Bracebridge Road
Sutton Coldfield
B74 2SL

Patents ADP number (if you know it)

8336646.0 01

If the applicant is a corporate body, give the country/state of its incorporation

4. Title of the invention

IMPROVED TARGETING DEVICE

5. Name of your agent (if you have one)

Barker Brettell

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

138 Hagley Road
Edgbaston
Birmingham
B16 9PW

Patents ADP number (if you know it)

7442494002

Country

Priority application number
(if you know it)Date of filing
(day/month/year)

6. Priority: Complete this section if you are declaring priority from one or more earlier patent applications, filed in the last 12 months.

7. Divisionals, etc: Complete this section only if this application is a divisional application or resulted from an entitlement dispute (see note f)

Number of earlier application

Date of filing
(day/month/year)8. Is a Patents Form 7/77 (Statement of inventorship and of right to grant of a patent) required in support of this request? Answer Yes' No
if:
a) any applicant named in part 3 is not an inventor, or
b) there is an inventor who is not named as an applicant,
or
c) any named applicant is a corporate body.
Otherwise answer NO (See note(d))

Patents Form 1/7

Patents Form 1/77

9. Enter the number of sheets for any of the following items you are filing with this form.
Do not count copies of the same document

Continuation sheets of this form -

Description 13 + 13

Claim(s) -

Abstract -

Drawing(s) 02 + 03

10. If you are also filing any of the following, state how many against each item.

Priority documents -

Translations of priority documents -

Statement of inventorship and right to grant of a patent (Patents Form 7/77) -

Request for preliminary examination -
(Patents Form 9/77)

Request for substantive examination -
(Patents Form 10/77)

Any other documents -
(please specify)

11. I/We request the grant of a patent on the basis of this application.

Signature *Barker Brettell*

Date

Barker Brettell

04.02.2004

12. Name and daytime telephone number of person to contact in the United Kingdom Ms. Lucy P. Truman Tel: 0121 456 1364

Warning

After an application for a patent has been filed, the Comptroller of the Patent Office will consider whether publication or communication of the invention should be prohibited or restricted under Section 22 of the Patents Act 1977. You will be informed if it is necessary to prohibit or restrict your invention in this way. Furthermore, if you live in the United Kingdom, Section 23 of the Patents Act 1977 stops you from applying for a patent abroad without first getting written permission from the Patent Office unless an application has been filed at least 6 weeks beforehand in the United Kingdom for a patent for the same invention and either no direction prohibiting publication or communication has been given, or any such direction has been revoked.

Notes

- If you need help to fill in this form or you have any questions, please contact the Patent Office on 01645 500505.
- Write your answers in capital letters using black ink or you may type them.
- If there is not enough space for all the relevant details on any part of this form, please continue on a separate sheet of paper and write "see continuation sheet" in the relevant part(s). Any continuation sheet should be attached to this form.
- If you have answered 'Yes' Patents Form 7/77 will need to be filed.
- Once you have filled in the form you must remember to sign and date it.
- For details of the fee and ways to pay please contact the Patent Office.

IMPROVED TARGETING DEVICE

The present invention relates to an improved targeting device, particularly for use in hip surgery and more particularly for use in hip replacement surgery or hip resurfacing surgery, and its use in surgery, particularly minimally invasive surgery.

When performing hip replacement surgery it is essential to be able to accurately determine where the centre of the osteotomised base of the femoral neck and the centreline through the femoral neck lie to allow correct reaming of the femoral neck and correct fitting of the replacement head onto the femoral neck.

The position of the femoral head does not always assist in determining the position of the centre of the osteotomised base of the femoral neck and the centreline through the femoral neck as the head may not be centrally positioned in relation to the femoral neck.

When performing hip resurfacing surgery it is essential to be able to accurately determine where the centreline through the femoral neck and femoral head lies to allow correct reaming of the femoral head and correct fitting of the prosthesis onto the femoral head.

The position of the femoral head does not always assist in determining the position of the centreline through the femoral neck and femoral head as the head may not be centrally positioned in relation to the femoral neck.

Accordingly there is the need for a targeting device which is suitable to allow the position of the centre of the femoral neck and/or the centreline through the femoral neck and femoral head (hereinafter reference to the centreline of the femoral neck is used to cover both of these references)

to be determined and that can be used in minimally invasive surgical methods.

The present invention provides in a first aspect a targeting device for use in minimally invasive hip surgery to allow the position of the centreline of the femoral neck to be located, which device comprises at least a first component suitable for location on an outer surface of the femoral neck and a second component suitable for marking the centreline of the femoral neck, wherein the first and second components are spaced apart from and parallel to one another and means is provided to alter the distance between the first and second components and means is provided to maintain the first and second components in a predetermined position relative to each other.

15 The present invention provides in a first embodiment a targeting device for use in minimally invasive hip surgery to allow the position of the centreline of the femoral neck to be located, which device comprises at least a first component suitable for receipt of a first guide wire and a second component suitable for receipt of a second guide wire, wherein the first and second components are spaced apart from and parallel to one another and means is provided to alter the distance between the first and second components and means is provided to maintain the first and second components in a predetermined position relative to each other.

20 25 The first and second components may be identical in configuration.

Preferably each of the first and second components is an elongate three dimensional shape, such as a cylinder. Each of the first and second components is preferably 2 to 10cm in length, more preferably 3 to 8cm and most preferably 4 to 6cm.

A central bore is preferably provided along the centre of each of the first and second components to receive a guide wire. The bore may have a diameter of 2 to 5mm, preferably 3 to 4mm, most preferably 3.3 to 3.5mm.

5

Each of the first and second components is most preferably cylindrical in shape with a central bore extending along its length, i.e. each of the first and second components is tubular.

10 The means provided to space the first and second components from each other and maintain the first and second components parallel to each other may be any suitable means.

15 Preferably the first component is provided with one or more runners, the or each runner is elongate and extends such that its longitudinal axis is perpendicular to that of the first component. Where there are two runners the runners are parallel to each other. The or each runner may be up to 10cm in length, preferably up to 8cm and most preferably up to 5cm.

20 The second component is preferably provided with means to movably, preferably slidably, engage the or each runner. The second component is positioned on the or each runner such that the longitudinal axis of the second component lies perpendicular to that of the or each runner and parallel to the longitudinal axis of the first component.

25

One or more springs may be provided between the first and second components to assist movement of the second component away from the first component, preferably the or each spring is provided around the or each runner.

30

The means to alter the distance between the first and second components may be any suitable means. The distance between the first and second components could be up to 10cm, preferably up to 8cm, more preferably up to 5cm.

5

The means to maintain the position of the second component relative to the first component may be any suitable means and is preferably the same means as the means to alter the distance between the first and second components.

10

Preferably there is provided a screw threaded bar received by corresponding screw threaded portions on each of the first and second components. The portions of the first and second components receiving the bar are preferably positioned on a common axis extending 15 perpendicular to and between the first and second components. The portions of first and second components receiving the bar are preferably protrusions having corresponding screw threaded apertures therethrough.

The screw threaded bar is preferably provided with a suitable means to 20 cause its rotation, for example a knob.

The first component is preferably mounted on an elongate support. The elongate support preferably has a handle portion distal from the first component. The first component is preferably mounted on the elongate 25 support perpendicular to the longitudinal axis of the elongate support.

In a most preferred embodiment the longitudinal axes of the or each runner and the screw threaded bar all lie parallel to the longitudinal axis of the elongate support and the longitudinal axes of the first and second 30 components each lie perpendicular to these axes and are spaced apart from and parallel to each other.

The device may comprise a first component and a second component only. Alternatively the device may further include at least a third component. The third component is preferably an "L" shaped bar having a first 5 elongate portion and a second elongate portion, the second elongate portion extending perpendicular to the first elongate portion. The length of the first elongate portion is greater than that of the second elongate portion.

10 The "L" shaped bar may be any suitable cross section but is preferably of circular, square or rectangular cross section. The "L" shaped bar may be solid or hollow.

15 The length of the first elongate portion may be from 10 to 30cm in length, preferably from 15 to 20cm.

20 The "L" shaped bar is positioned parallel to and spaced apart from the first component. The "L" shaped bar is preferably mounted on the elongate support on the side of the first component opposite to that of the second component.

25 The provision of the "L" shaped bar is particularly advantageous in hip resurfacing surgery where it ensures that the first component is in line with the outer surface of the femoral neck. Without the provision of the "L" shaped bar this cannot be done accurately as the femoral head is in the way and it is not therefore possible to see whether the first component is actually positioned in line with the outer surface of the femoral neck. In hip replacement surgery the provision of the "L" shaped bar provides additional assistance in locating and steadyng the targeting device.

The "L" shaped bar is mounted on the elongate support such that the first elongate portion is parallel to the first component and the "L" shaped bar it is movable along the longitudinal axis of the first elongate portion.

5 The "L" shaped bar being movable along the longitudinal axis of the first portion allows for its use with different sized hip joints by allowing for the device to be used around different sized femoral heads.

10 The "L" shaped bar is mounted on the elongate support such that the free end of the second elongate portion extends as far as the longitudinal axis through the first component. The free end of the second elongate portion may be provided with a plate perpendicular thereto which lies along the longitudinal axis through the first component. Accordingly the length of the second elongate portion, including any plate, is the same as the 15 distance between the first component and the first elongate portion of the "L" shaped bar, this distance is preferably 2 to 5cm, most preferably 2 to 3cm.

20 The present invention provides in a second embodiment a targeting device for use in minimally invasive hip surgery to allow the position of the centreline of the femoral neck to be located, wherein the first component is an "L" shaped bar according to the first embodiment and the second component is a second component according to the first embodiment.

25 Preferably the second component is movable in relation to the "L" shaped bar such that the second component is spaced apart from the free end of the second elongate portion of the "L" shaped bar by a distance that is half of the diameter of the femoral neck to allow the centreline through the femoral neck to be determined.

The targeting device may be manufactured from any suitable material, for example metal such as stainless steel or surgical steel or a plastics material.

5 In a second aspect the targeting device of the present invention may be used in a method of locating the centre of the osteotomised base of the femoral neck, which method comprises:

-measuring the diameter of the femoral neck;

10

-dividing the diameter of the femoral neck by two to give value X and setting the distance between the first and second components of the targeting device as X;

15 -determining the mid line through the femoral neck in the AP plane and running a guide wire along this line through the first component of the targeting device of the present invention;

20 -determining the centre of the osteotomised femoral neck by rotation of the second component around the guide wire running through the first component.

25 Preferably a second guide wire is passed through the second component to mark the centre of the osteotomised femoral neck. The targeting device may then be removed leaving one or both of the guide wires in place.

Preferably the diameter of the femoral neck is measured using callipers.

30 The "L" shaped bar may be provided and positioned such that the free end of the second elongate portion of the "L" shaped bar rests on the

femoral neck on the midline in the AP plane. This assists in location and use of the targeting device.

In a third aspect the targeting device of the present invention may be used
5 in a method of locating the centreline through the femoral neck and femoral head, which method comprises:

- measuring the diameter of the femoral neck;
- 10 -dividing the diameter of the femoral neck by two to give value X and setting the distance between the first and second components of the targeting device as X;
- determining the mid line through the femoral neck in the AP
15 plane;
- positioning the free end of the second elongate portion of the "L" shaped bar on the femoral neck on the midline in the AP plane;
- 20 -determining the centreline through the femoral neck and femoral head by position of the second component.

A guide wire may be passed through the first component once its position has been determined using the "L" shaped bar.

25 Preferably a second guide wire is passed through the second component to mark the central longitudinal axis of the femoral neck. The targeting device may then be removed leaving one or both of the guide wires in place.

30 Preferably the diameter of the femoral neck is measured using callipers.

A specific embodiment of the present invention will now be described, by means of example only, with reference to the drawings, in which:

5 Figure 1 shows, in plan view, a schematic drawing of a first targeting device according to the present invention; and

Figure 2 shows, in plan view, a schematic drawing of a further targeting device according to the present invention.

10

Figure 1 shows a targeting device 1 suitable for use in minimally invasive hip replacement surgery. The device 1 comprises an elongate support 2 having a handle portion 3 at one end having a central longitudinal axis B. Secured to the end of the elongate support distal from the handle portion 15 3, by welding, is a first component 4 suitable for receiving a guide wire.

The first component 4 comprises a tube having a central longitudinal axis A lying perpendicular to the longitudinal axis B of the elongate support 2.

20 Two elongate runners 5, 6 are provided extending tangentially to the first component 4. The runners 5, 6 are parallel to and spaced apart from each other and their central longitudinal axes C1, C2 run parallel to axis B of the elongate support 2. The runners 5, 6 are secured to the first component 4 by welding.

25

A second component 7 is slidably mounted on the runners 5, 6. The second component 7 comprises a tube having a central longitudinal axis D parallel to axis A of the first component 4.

30 The second component 7 is provided with two suitably sized shoulders 8a, 8b having apertures 9a, 9b (not shown) therein to receive the runners 5,

6. The shoulders 8a, 8b are positioned such that the runners form an identical tangent to the second component 7 as they do to the first component 4.

5 Each of the first 4 and second 7 components is provided with an integral protrusion 10a, 10b positioned diametrically opposite to the runners 5, 6 and having a screw threaded aperture therethrough. A correspondingly screw threaded bar 11 is provided extending through the apertures in the protrusions 10a, 10b and rotation of this bar 11 causes movement of the 10 second component 7 relative to the first component 4 along the runners 5, 6. The bar 11 is provided with a knob 12 to assist in its manual rotation.

15 Springs 13a, 13b are provided around the runners 5, 6 between the first 4 and second 7 components to assist movement of the second component 7 away from the first component 4.

In use the targeting device of the present invention assists in locating the central longitudinal axis of the femoral neck. In minimally invasive hip replacement surgery as described in International Patent Publication No 20 WO 03/065906 of the same inventor the femoral neck osteotomy is performed and the osteomised femur is exposed as described. To locate the centre of the osteomised femoral neck, in the AP and lateral plane, the diameter of the femoral neck is measured with callipers and the radius is calculated by dividing the diameter by two.

25 The distance of the centre of the second component 7 from the centre of the first component 4 is adjusted by rotation of the screw threaded bar 11 until the distance is the same as the measured diameter divided by two.

30 The centre line of the femoral neck in the AP plane is determined and a first guide wire is inserted in line with this centre line. The first guide

wire is passed through the first component 4 of the targeting device 1. The second component 7 of the targeting device is then used to locate the centre in the AP and lateral plane by rotation around the first guide wire. The centre is then marked with a second guide wire and the targeting 5 device can be removed leaving one or both of the guide wires in place.

The femoral neck can then be reamed accurately and centrally and the resultant prosthesis will be accurately and centrally positioned.

- 10 Figure 2 shows a further targeting device 100 suitable for use in minimally invasive hip surgery, especially minimally invasive hip resurfacing surgery. In Figure 2 the targeting device 100 is similar to targeting device 1 and like numbers relate to like parts.
- 15 Targeting device 100 is additionally provided with an "L" shaped bar 101 comprising a first elongate portion 102 and a second elongate portion 103 where the first elongate portion is of greater length than the second elongate portion.
- 20 The "L" shaped bar 101 is mounted on the elongate support 2 such that the first elongate portion 102 is parallel to the first component 4. The second elongate portion 103 is perpendicular to the first elongate portion 102 and the free end of the second portion is provided with a plate 104 that lies parallel with the first component 4.
- 25 The "L" shaped bar 101 is sized and mounted on the elongate support 2 such that the plate 104 lies along the longitudinal axis through the first component 4.
- 30 The "L" shaped bar 101 is movable along the longitudinal axis through the first elongate portion 102. The "L" shaped bar 101 is mounted on the

elongate support 2 by means of a channel (not shown) provided through a block 105 secured by welding to the elongate support 2 in which the first elongate portion 102 is received. A bolt 106 is provided extending into the channel through the block 105 to contact the first elongate portion and 5 lock the "L" shaped bar in a desired position.

In use the targeting device of the present invention assists in locating the central longitudinal axis of the femoral neck. In minimally invasive hip resurfacing surgery as described in International Patent Publication No

10 WO 03/065906 of the same inventor the femoral head can be delivered into the wound at 90° of internal rotation, 45° of flexion and as much adduction as possible as described. To locate the centreline through the femoral neck and femoral head, in the AP and lateral plane, the diameter of the femoral neck is measured with callipers and the radius is calculated 15 by dividing the diameter by two.

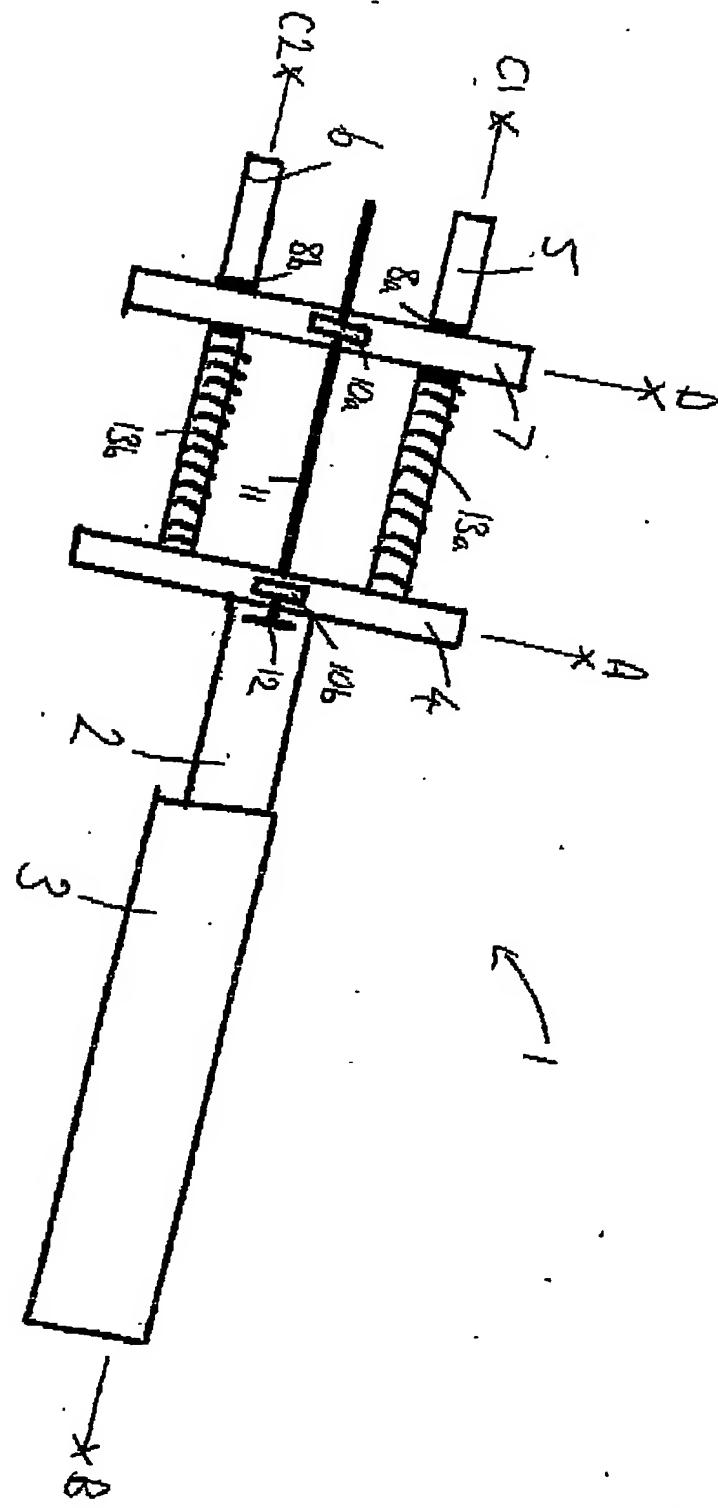
The distance of the centre of the second component 7 from the centre of the first component 4 is adjusted by rotation of the screw threaded bar 11 until the distance is the same as the measured diameter divided by two.

20 The centre line of the femoral neck in the AP plane is determined and the plate 104 of the "L" shaped bar 101 is positioned on the centreline in the AP plane. A first guide wire may be passed through the first component 4 of the targeting device 1 and this is certain to be in line with the 25 centreline in the AP plane owing to the use of the "L" shaped bar. The second component 7 of the targeting device is used to locate the centre in the AP and lateral plane by rotation around the first guide wire and/or the "L" shaped bar. The centre is then marked with a guide wire passed through the second component and the targeting device can be removed 30 leaving the guide wire in place.

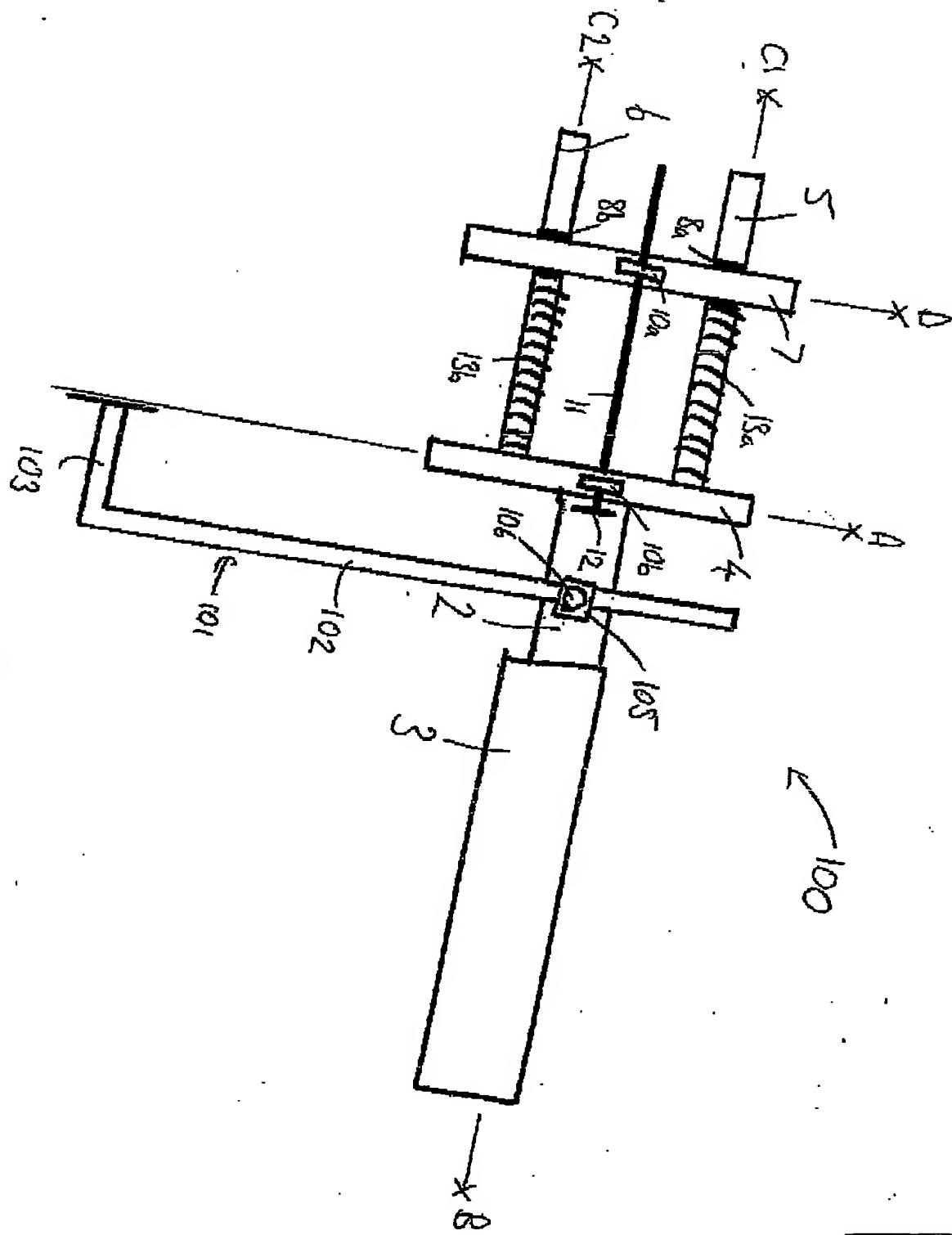
13

The femoral head can then be reamed accurately and centrally and the resultant prosthesis will be accurately and centrally positioned.

1/2



2/a



IMPROVED TARGETING DEVICE

The present invention relates to an improved targeting device, particularly for use in hip surgery and more particularly for use in hip replacement surgery or hip resurfacing surgery, and its use in surgery, particularly minimally invasive surgery.

When performing hip replacement surgery it is essential to be able to accurately determine where the centre of the base of the femoral neck and 10 the centreline through the femoral neck lie to allow correct reaming fitting of the replacement head.

The position of the femoral head does not always assist in determining the position of the centre of the femoral neck and the centreline through the 15 femoral neck as the head may not be centrally positioned in relation to the femoral neck due to deformity of the femoral head.

When performing hip resurfacing surgery it is essential to be able to accurately determine where the centreline through the femoral neck and 20 femoral head lies to allow correct reaming of the femoral head and correct fitting of the prosthesis onto the femoral head.

The position of the femoral head does not always assist in determining the position of the centreline through the femoral neck and femoral head as 25 the head may not be centrally positioned in relation to the femoral neck due to deformity of the femoral head.

Accordingly there is the need for a targeting device which is suitable to allow the position of the centre of the femoral neck and/or the centreline 30 through the femoral neck and femoral head (hereinafter reference to "the centreline of the femoral neck" is used to cover both of these references)

to be determined, which device can be used in minimally invasive surgical methods and in conventional open surgical methods.

The present invention provides in a first aspect a targeting device for use in open, or minimally invasive, hip surgery to allow the position of the centreline of the femoral neck to be located, which device comprises at least a first component having a portion suitable for location on an outer surface of the femoral neck and a second component having a portion suitable for marking the centreline of the femoral neck, wherein the first and second components are spaced apart from and parallel to one another and means is provided to alter the distance between the first and second components and means is provided to maintain the first and second components in a predetermined position relative to each other.

10 The present invention provides in a first embodiment a targeting device for use in open, or minimally invasive, hip surgery to allow the position of the centreline of the femoral neck to be located, which device comprises at least a first component having a portion suitable for receipt of a first guide wire and a second component having a portion suitable for receipt of a second guide wire, wherein the first and second components are spaced apart from and parallel to one another and means is provided to alter the distance between the first and second components and means is provided to maintain the first and second components in a predetermined position relative to each other.

15

20

25

The first and second components may be identical in configuration.

Preferably each of the first and second components includes an elongate three dimensional shape, such as a cylinder, for location on an outer surface of the femoral neck or for marking the centreline of the femoral neck. Each three dimensional shape of the first and second components is

30

preferably 2 to 8cm in length, more preferably 3 to 6cm in length and most preferably 4 to 5cm in length.

5 A central bore is preferably provided along the centre of each of the elongate three dimensional shapes of the first and second components to receive a guide wire. The bore may have a diameter of 1.5 to 5mm, preferably 3 to 4mm, most preferably 3.2 to 3.5mm.

10 Each of the elongate three dimensional shapes of the first and second components is most preferably cylindrical in shape with a central bore extending along its length, i.e. each of the three dimensional elongate shapes of the first and second components is tubular.

15 The means provided to space the first and second components from each other and maintain the first and second components parallel to each other may be any suitable means.

20 Preferably the first component is provided with one or more runners, the or each runner is elongate and extends such that its longitudinal axis is perpendicular to that of the first component. The or each runner is preferably secured, for example by welding, to the first component. Where there are two or more runners the runners are parallel to each other. The or each runner may be up to 6cm in length, preferably up to 5cm and most preferably up to 3 or 2cm.

25

30 The second component is preferably provided with means to movably, preferably slidably, engage the or each runner. The second component is positioned on the or each runner such that the longitudinal axis of the second component lies perpendicular to that of the or each runner and parallel to the longitudinal axis of the first component. In use the second

component is movable along the or each runner to alter the distance between the first and second component.

Alternatively the second component is provided with one or more runners, the or each runner is elongate and extends such that its longitudinal axis is perpendicular to that of the second component. The or each runner is preferably secured, for example by welding, to the second component. Where there are two or more runners the runners are parallel to each other. The or each runner may be up to 6cm in length, preferably up to 5cm and most preferably up to 3 or 2cm.

The first component is preferably provided with means to movably, preferably slidably, engage the or each runner. The first component is positioned on the or each runner such that the longitudinal axis of the first component lies perpendicular to that of the or each runner and parallel to the longitudinal axis of the first component. In use the second component is movable in relation to the first component by the or each runner moving in relation to the first component. This arrangement ensures that the device is compact and when the first and second components are close together the or each runner extend out of the incision to avoid tissue damage.

One or more springs may be provided between the first and second components to assist movement of the second component away from the first component, preferably the or each spring is provided around the or each runner.

In a most preferred embodiment each of the first and second components is provided with a mounting portion and an elongate three dimensional shaped portion. The mounting portion is suitably sized and shaped to receive or co-operate with the elongate three dimensional shaped portion,

the or each runner and the means provided to space the first and second components from each other and maintain the first and second components parallel to each other. The elongate three dimensional shaped portions are preferably releasably secured to the mounting portions, most 5 preferably each of the elongate three dimensional shaped portions is tubular and has an external screw threaded portion. The mounting portions preferably each have one or more apertures with corresponding internal screw threaded portions to receive and hold a tubular portion. This allows the tubular portions to be removed for cleaning or to be 10 disposed of and replaced.

The targeting device may also be provided with a repositioning device. The repositioning device preferably comprises a plate to be secured to the targeting device and having one, two or more tubular components, sized 15 to receive a guide wire, extending therefrom. The plate is most preferably received by the apertures in the first and second components when the first and second components are positioned as close together as possible. This fixing is quick and stable.

20 The means to alter the distance between the first and second components may be any suitable means. The distance between the first and second components could be up to 3cm, preferably up to 2.5cm.

25 The means to maintain the position of the second component relative to the first component may be any suitable means and is preferably the same means as the means to alter the distance between the first and second components.

30 Preferably there is provided a screw threaded bar received by corresponding screw threaded portions on each of the first and second components. The portions of the first and second components receiving

the bar are preferably positioned on a common axis extending perpendicular to and between the first and second components. The portions of first and second components receiving the bar are preferably protrusions having corresponding screw threaded apertures therethrough.

5

The screw threaded bar is preferably provided with a suitable means to cause its rotation, for example a knob.

10 In a most preferred embodiment the longitudinal axes of the or each runner and the screw threaded bar all lie parallel to the longitudinal axis of the elongate support and the longitudinal axes of the first and second components each lie perpendicular to these axes and are spaced apart from and parallel to each other.

15 The first component is preferably mounted on an elongate support. The elongate support preferably has a handle portion distal from the first component. ~~The first component is preferably mounted on the elongate support perpendicular to the longitudinal axis of the elongate support.~~

20 Alternatively the first component is mounted on an elongate support and the second component is mounted on a bar telescopically received within the elongate support. The bar is preferably slidably moved into and out of the elongate support by means of a screw thread system.

25 Preferably a rotatable shaft is provided within the elongate support having a screw thread corresponding to a screw thread provided on the telescopically received bar. The screw thread on the bar may be external and the screw thread on the shaft may be internal or vice versa. The rotatable shaft is preferably provided with a knob to assist in its rotation.

30 Rotation of the rotatable shaft causes movement of the telescopically received bar into and out of the elongate support. When rotation of the

rotatable shaft is stopped the first and second components are held in the desired position relative to each other.

5 The screw thread system is preferably provided with a suitable means to cause the bar to move into and out of the elongate support by rotation, for example a knob.

10 In a most preferred embodiment the longitudinal axes of the or each runner the elongate support and the telescopically received bar all lie parallel to the longitudinal axis of the elongate support and the longitudinal axes of the first and second components each lie perpendicular to these axes and are spaced apart from and parallel to each other.

15 The device may comprise a first component and a second component only. Alternatively the device may further include at least a third component. The third component is preferably an "L" shaped bar having a first elongate portion and a second elongate portion, the second elongate portion extending generally perpendicularly to the first elongate portion.
20 The length of the first elongate portion is greater than that of the second elongate portion.

25 The "L" shaped bar may be any suitable cross section but is preferably of circular, square or rectangular cross section, the "L" shaped bar is preferably of a cross section that prevents its rotation around its longitudinal axis to ensure that a desired alignment is maintained. The "L" shaped bar may be solid or hollow.

30 The length of the first elongate portion may be from 10 to 30cm in length, preferably from 15 to 25cm and most preferably from 15 to 20cm.

The "L" shaped bar is positioned parallel to and spaced apart from the first component. The "L" shaped bar is preferably mounted on the elongate support on the side of the first component opposite to that of the second component.

5

The provision of the "L" shaped bar is particularly advantageous in hip resurfacing surgery where it ensures that the first component is in line with the outer surface of the femoral neck. Without the provision of the "L" shaped bar this cannot be done accurately as the femoral head is in the way and it is not therefore possible to see whether the first component is actually positioned in line with the outer surface of the femoral neck. In hip replacement surgery the provision of the "L" shaped bar provides additional assistance in locating and steadyng the targeting device.

15 The "L" shaped bar is mounted on the elongate support such that the first elongate portion is parallel to the first component and the "L" shaped bar and the "L"-shaped bar is movable along the longitudinal axis of the first elongate portion.

20 The "L" shaped bar being movable along the longitudinal axis of the first portion allows for its use with different sized hip joints by allowing for the device to be used around different sized femoral heads.

25 The "L" shaped bar is mounted on the elongate support such that the free end of the second elongate portion extends as far as the longitudinal axis through the first component. The free end of the second elongate portion may be provided with a plate perpendicular thereto which lies along the longitudinal axis through the first component. Accordingly the length of the second elongate portion, including any plate, is the same as the 30 distance between the first component and the first elongate portion of the

"L" shaped bar, this distance is preferably 2 to 5cm, most preferably 2 to 3cm.

The present invention provides in a second embodiment a targeting device
5 for use in open, or minimally invasive, hip surgery to allow the position
of the centreline of the femoral neck to be located, wherein the first
component is an "L" shaped bar according to the first embodiment and
the second component is a second component according to the first
embodiment.

10

Preferably the second component is movable in relation to the "L" shaped
bar such that the second component is spaced apart from the free end of
the second elongate portion of the "L" shaped bar by a distance that is
half of the diameter of the femoral neck to allow the centreline through
15 the femoral neck to be determined.

The targeting device may be manufactured from any suitable material, for
example metal such as stainless steel or surgical steel or a plastics
material.

20

In a second aspect the targeting device of the present invention may be
used in a method of locating the centre of the osteotomised base of the
femoral neck at the head-neck junction, which method comprises:

25

-measuring the diameter of the femoral neck;

-dividing the diameter of the femoral neck by two to give value X
and setting the distance between the first and second components of
the targeting device as X;

30

-determining the mid line through the femoral neck in the AP plane and running a guide wire along this line through the first component of the targeting device of the present invention.

5 Preferably a second guide wire is passed through the second component to mark the centre of the osteotomised femoral neck. The targeting device may then be removed leaving at least the second guide wire in place.

10 Preferably the diameter of the femoral neck is measured using callipers. Alternatively the diameter may be measured using a suitable calibrated gauge.

15 The "L" shaped bar may be provided and positioned such that the free end of the second elongate portion of the "L" shaped bar rests on the femoral neck on the midline in the AP plane. This assists in location and use of the targeting device.

20 In a third aspect the targeting device of the present invention may be used in a method of locating the centreline through the femoral neck and femoral head, which method comprises:

25 -measuring the diameter of the femoral neck;

-dividing the diameter of the femoral neck by two to give value X and setting the distance between the first and second components of the targeting device as X;

-determining the mid line through the femoral neck in the AP plane;

- positioning the free end of the second elongate portion of the "L" shaped bar on the femoral neck on the midline in the AP plane;
- 5 -determining the centreline through the femoral neck and femoral head by position of the second component.

A guide wire may be passed through the first component once its position has been determined using the "L" shaped bar.

- 10 Preferably a second guide wire is passed through the second component to mark the central longitudinal axis of the femoral neck. The targeting device may then be removed leaving at least the second guide wire in place.
- 15 Preferably the diameter of the femoral neck is measured using callipers. Alternatively the diameter may be measured using a suitable calibrated gauge.

- 20 The present invention further provides a gauge for measuring the diameter of a femoral neck, which gauge comprises a first component and a second component wherein at least one of the first and second components is movable relative to the other said component and means is provided to determine the distance between the first and second components.

- 25 Preferably the first component is fixed in position and the second component is movable in relation to the first component or vice versa.
- 30 Preferably the device is provided with a handle portion to which the first component is fixedly mounted, preferably by means of a generally semi-circular joining member.

The second component is preferably slidably mounted on or within the handle portion and is movable towards and away from the second component.

- 5 The distance between the first and second component may be determined by any suitable means, for example by the provision of a scale to show the distance between the first and second component depending on the relative positions of the first and second components.
- 10 A specific embodiment of the present invention will now be described, by means of example only, with reference to the drawings, in which:

15 **Figure 1a, 1b and 1c** show, in plan view, schematic drawings of a first, second and third targeting device according to the present invention;

Figure 2 shows, in plan view, a schematic drawing of a further targeting device according to the present invention; and

20 **Figure 3** shows a cross section through a gauge for measuring the diameter of a femoral neck.

25 Figures 1a and 1b shows a targeting device 1 suitable for use in open, or minimally invasive, hip replacement surgery. The device 1 comprises an elongate support 2 having a handle portion 3 at one end having a central longitudinal axis B. Secured to the end of the elongate support distal from the handle portion 3, by welding, is a first component 4 suitable for receiving a guide wire.

In a first embodiment, shown in Figure 1a, the first component 4 comprises a tube having a central longitudinal axis A lying perpendicular to the longitudinal axis B of the elongate support 2.

5 Two elongate runners 5, 6 are provided extending tangentially to the first component 4. The runners 5, 6 are parallel to and spaced apart from each other and their central longitudinal axes C1, C2 run parallel to axis B of the elongate support 2. The runners 5, 6 are secured to the first component 4 by welding.

10

A second component 7 is slidably mounted on the runners 5, 6. The second component 7 comprises a tube having a central longitudinal axis D parallel to axis A of the first component 4.

15 The second component 7 is provided with two suitably sized shoulders 8a, 8b having apertures 9a, 9b (not shown) therein to receive the runners 5, 6. The shoulders 8a, 8b are positioned such that the runners form an identical tangent to the second component 7 as they do to the first component 4.

20

Each of the first 4 and second 7 components is provided with an integral protrusion 10a, 10b positioned diametrically opposite to the runners 5, 6 and having a screw threaded aperture therethrough. A correspondingly screw threaded bar 11 is provided extending through the apertures in the protrusions 10a, 10b and rotation of this bar 11 causes movement of the second component 7 relative to the first component 4 along the runners 5, 6. The bar 11 is provided with a knob 12 to assist in its manual rotation.

30 In a second embodiment, shown in Figure 1b, the first component 4 comprises a tube having a central longitudinal axis A lying perpendicular to the longitudinal axis B of the elongate support 2. A second component

7 is provided and comprises a tube having a central longitudinal axis D parallel to axis A of the first component 4.

Two elongate runners 5, 6 are provided extending tangentially to the first
5 component 4. The runners 5, 6 are parallel to and spaced apart from each other and their central longitudinal axes C1, C2 run parallel to axis B of the elongate support 2. The runners 5, 6 are secured to the second component 7 by welding.

10 The first component 4 slidably receives the runners 5, 6 by provision of with two suitably sized shoulders 8a, 8b having apertures 9a, 9b (not shown) therein to receive the runners 5, 6. The shoulders 8a, 8b are positioned such that the runners form an identical tangent to the first component 4 as they do to the second component 7.

15

Each of the first 4 and second 7 components is provided with an integral protrusion-10a,-10b-positioned-diametrically opposite to the runners 5, 6 and having a screw threaded aperture therethrough. A correspondingly screw threaded bar 11 is provided extending through the apertures in the

20 protrusions 10a, 10b and rotation of this bar 11 causes movement of the second component 7 relative to the first component 4 by the runners 5, 6 sliding through the apertures 9a and 9b in shoulders 8a, 8b. The bar 11 is provided with a knob 12 to assist in its manual rotation.

25 In both embodiments springs 13a, 13b are provided around the runners 5, 6 between the first 4 and second 7 components to assist movement of the second component 7 away from the first component 4.

30 In a third embodiment, shown in Figure 1c, the first component 4 includes a mounting portion 4a and a tube 4b having a central longitudinal axis A lying perpendicular to the longitudinal axis B of the elongate

support 2. A second component 7 is provided and includes a mounting portion 7a and a tube 7b having a central longitudinal axis D parallel to axis A of the first component 4. The tubes 4b and 7b have external screw threaded portions (not shown) and are releasably secured to the mounting portions 4a and 7a by the provision of apertures in the mounting portions with internal screw threaded portions (not shown) corresponding to external screw threads of the tubes 4b, 7b. The tubes 4b, 7b can then be removed for cleaning or can be disposed of and replaced.

5 The mounting portion 4a of the first component 4 is secured to the elongate support 2 by welding. The mounting portion 7a of the second component 7 is secured to a bar 16 by welding. The bar 16 is telescopically received by the elongate support 2 which has a bore along its length.

10 15 Two elongate runners 5, 6 are provided extending from the mounting portion 7a of the second component 7. The runners 5, 6 are parallel to and spaced apart from each other and their central longitudinal axes C1, C2 run parallel to axis B of the elongate support 2. The runners 5, 6 are secured to the second component 7 by welding.

20 The mounting portion 4a of the first component 4 slidably receives the runners 5, 6 by provision of two suitably sized apertures 19a, 19b therein to receive the runners 5, 6.

25 The second component 7 is moved relative to the first component 4 by the bar 16 telescopically moving into and out of the bore in the elongate support 2. During this movement the runners 5, 6 hold the position of the second component and prevent the bar 16 from rotating. To cause 30 movement of the bar 16 along the bore of the elongate support 2 the bar 16 is provided with an external screw thread 17 around its end positioned

farthest from the second component. The elongate support 2 is provided with a rotatable shaft 18 positioned within the elongate support and having an internal screw thread (not shown) corresponding to the external screw thread of the bar 16. The rotatable shaft is provided with a knob 5 21 at the end thereof, and positioned on the outside of the elongate support 2, to allow rotation of the shaft 18. The rotation of the shaft 18 causes movement of the bar 16 by means of the action of the corresponding screw threads and the prevention of rotation of the bar 16 by means of the runners 5, 6.

10

In use the targeting devices of the present invention assist in locating the central longitudinal axis of the femoral neck. In minimally invasive hip replacement surgery as described in International Patent Publication No WO 03/065906 of the same inventor, or in open hip replacement surgery, 15 the diameter of the femoral neck is measured and the radius is calculated by dividing the diameter by two.

The distance of the centre of the second component 7 from the centre of the first component 4 is adjusted by rotation of the screw threaded bar 11 20 until the distance is the same as the measured diameter divided by two.

The centre line of the femoral neck in the AP plane is determined and a first guide wire is inserted in line with this centre line. The first guide wire is passed through the first component 4 of the targeting device 1. 25 The second component 7 of the targeting device is then used to locate the centre in the AP and lateral plane by rotation around the first guide wire. The centre is then marked with a second guide wire and the targeting device can be removed leaving the second guide wire in place.

30 The femoral neck can then be reamed accurately and centrally and the resultant prosthesis will be accurately and centrally positioned.

Figure 2 shows a further targeting device 100 suitable for use in open, or minimally invasive, hip surgery, especially minimally invasive hip resurfacing surgery. In Figure 2 the targeting device 100 is similar to targeting device 1 of the embodiment of Figure 1a and like numbers relate 5 to like parts.

Targeting device 100 is additionally provided with an "L" shaped bar 101 comprising a first elongate portion 102 and a second elongate portion 103 where the first elongate portion is of greater length than the second 10 elongate portion.

The "L" shaped bar 101 is mounted on the elongate support 2 such that the first elongate portion 102 is parallel to the first component 4. The second elongate portion 103 is perpendicular to the first elongate portion 15 102 and the free end of the second portion is provided with a plate 104 that lies parallel with the first component 4.

The "L" shaped bar 101 is sized and mounted on the elongate support 2 such that the plate 104 lies along the longitudinal axis through the first 20 component 4.

The "L" shaped bar 101 is movable along the longitudinal axis through the first elongate portion 102. The "L" shaped bar 101 is mounted on the elongate support 2 by means of a channel (not shown) provided through a block 105 secured by welding to the elongate support 2 in which the first 25 elongate portion 102 is received. A bolt 106 is provided extending into the channel through the block 105 to contact the first elongate portion and lock the "L" shaped bar in a desired position.

30 The "L" shaped bar can also be used with the targeting device of Figures 1b and 1c in the same manner as described in relation to Figure 1a

In use the targeting device of the present invention assists in locating the central longitudinal axis of the femoral neck. In minimally invasive hip resurfacing surgery as described in International Patent Publication No WO 03/065906 of the same inventor the femoral head can be delivered

5 into the wound at 90° of internal rotation, 45° of flexion and as much adduction as possible as described. To locate the centreline through the femoral neck and femoral head, in the AP and lateral plane, in open, or minimally invasive, hip resurfacing surgery the diameter of the femoral neck is measured and the radius is calculated by dividing the diameter by

10 two.

The distance of the centre of the second component 7 from the centre of the first component 4 is adjusted by rotation of the screw threaded bar 11 until the distance is the same as the measured diameter divided by two.

15 The centre line of the femoral neck in the AP plane is determined and the plate-104-of-the—"L"-shaped-bar-101-is positioned on the centreline in the AP plane. A first guide wire may be passed through the first component 4 of the targeting device 1 and this is certain to be in line with the centreline in the AP plane owing to the use of the "L" shaped bar. The second component 7 of the targeting device is used to locate the centre in the AP and lateral plane by rotation around the first guide wire and/or the "L" shaped bar. The centre is then marked with a guide wire passed through the second component and the targeting device can be removed

20

25 leaving the second guide wire in place.

The femoral head can then be reamed accurately and centrally and the resultant prosthesis will be accurately and centrally positioned.

30 Figure 3 shows a gauge for use in measuring the diameter of a femoral head or neck. The gauge 29 comprises an elongate handle 30 having a

first component 31 fixedly secured thereto such that the centre of the first component lies on the longitudinal axis through the handle 30. The first component is fixed to the handle by means of a semi-circular joining member 32 extending between the handle 30 and the first component 31.

5 The semi-circular joining member 32 allows the device to be used around a generally cylindrical bone to determine the diameter thereof.

A second component 33 is provided slidably attached within a channel 36 in the handle 30 by a rod 37 and seal 38 arrangement. The second component also lies on the longitudinal axis through the handle and is positioned between the handle 30 and the first component 31. A measuring scale 34 is provided (shown in dotted lines), calibrated to show the distance between the first and second components as the second component moves in relation to the first component. A marker 35 (shown in dotted lines) secured to the second component 33 moves along the scale as the second component 33 moves in relation to the first component 31 showing the distance between the two components.

The device is sized such that the maximum distance between the two components is 4cm.

1/5

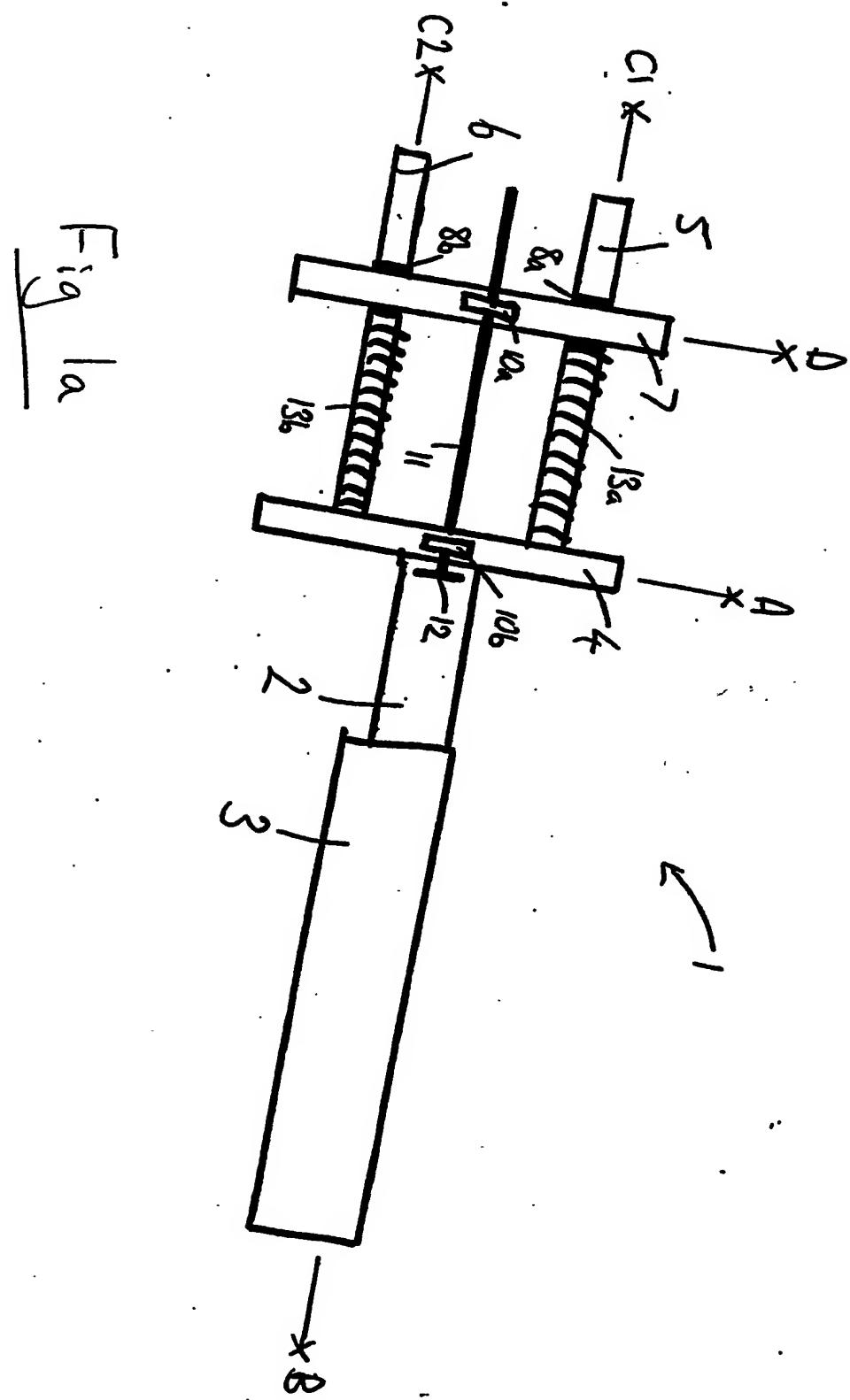


Fig 1a

2/5

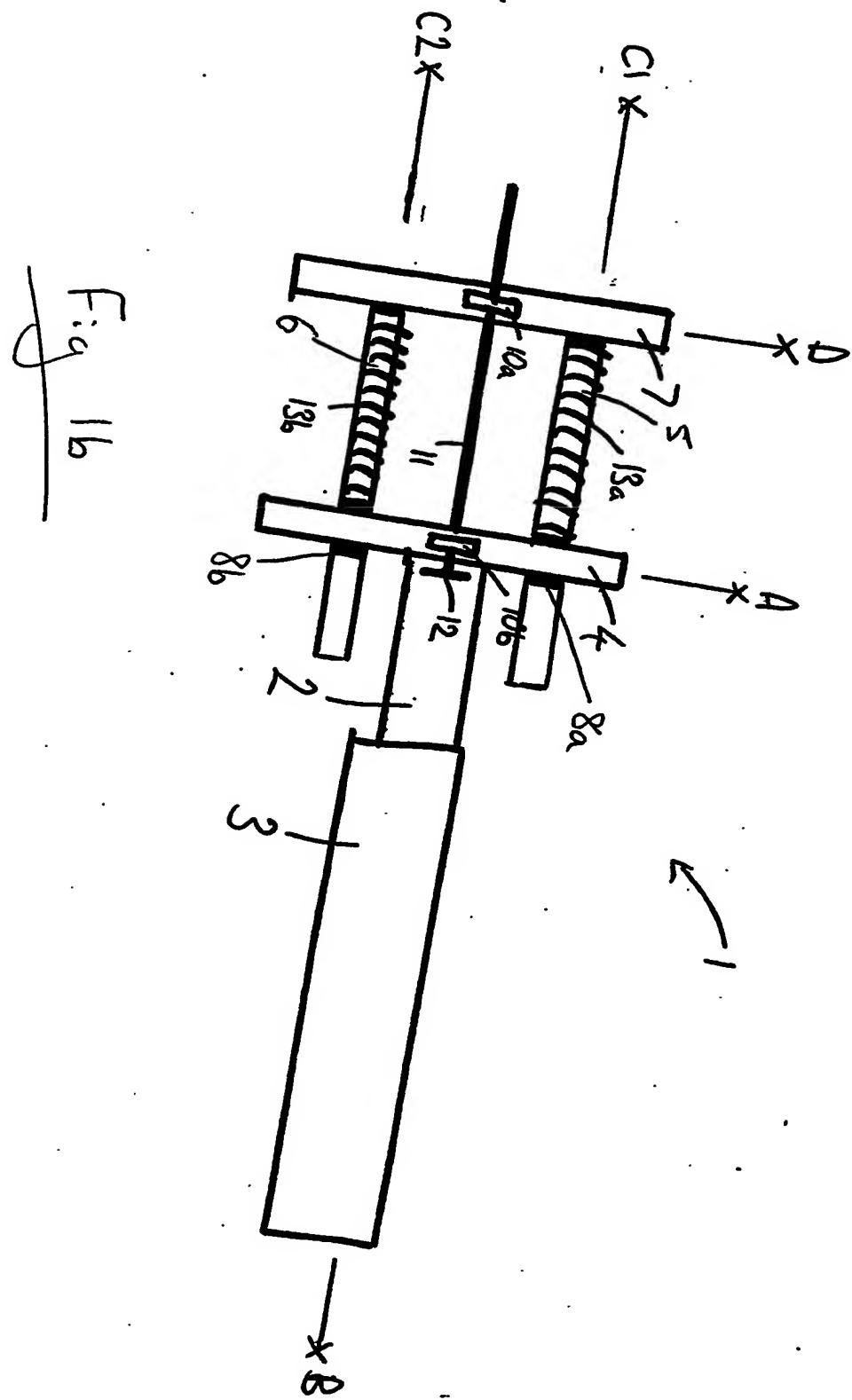
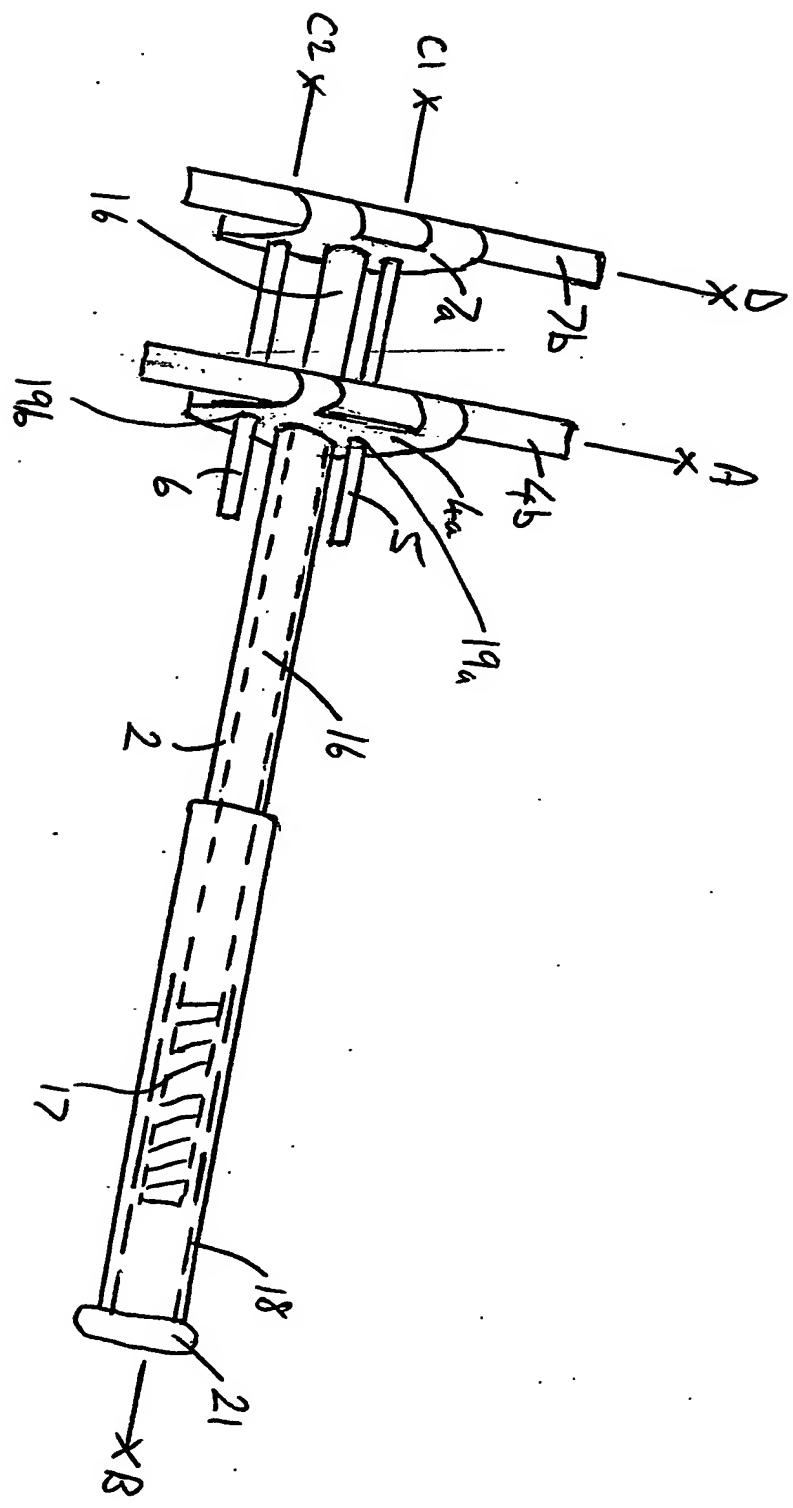


Fig 16

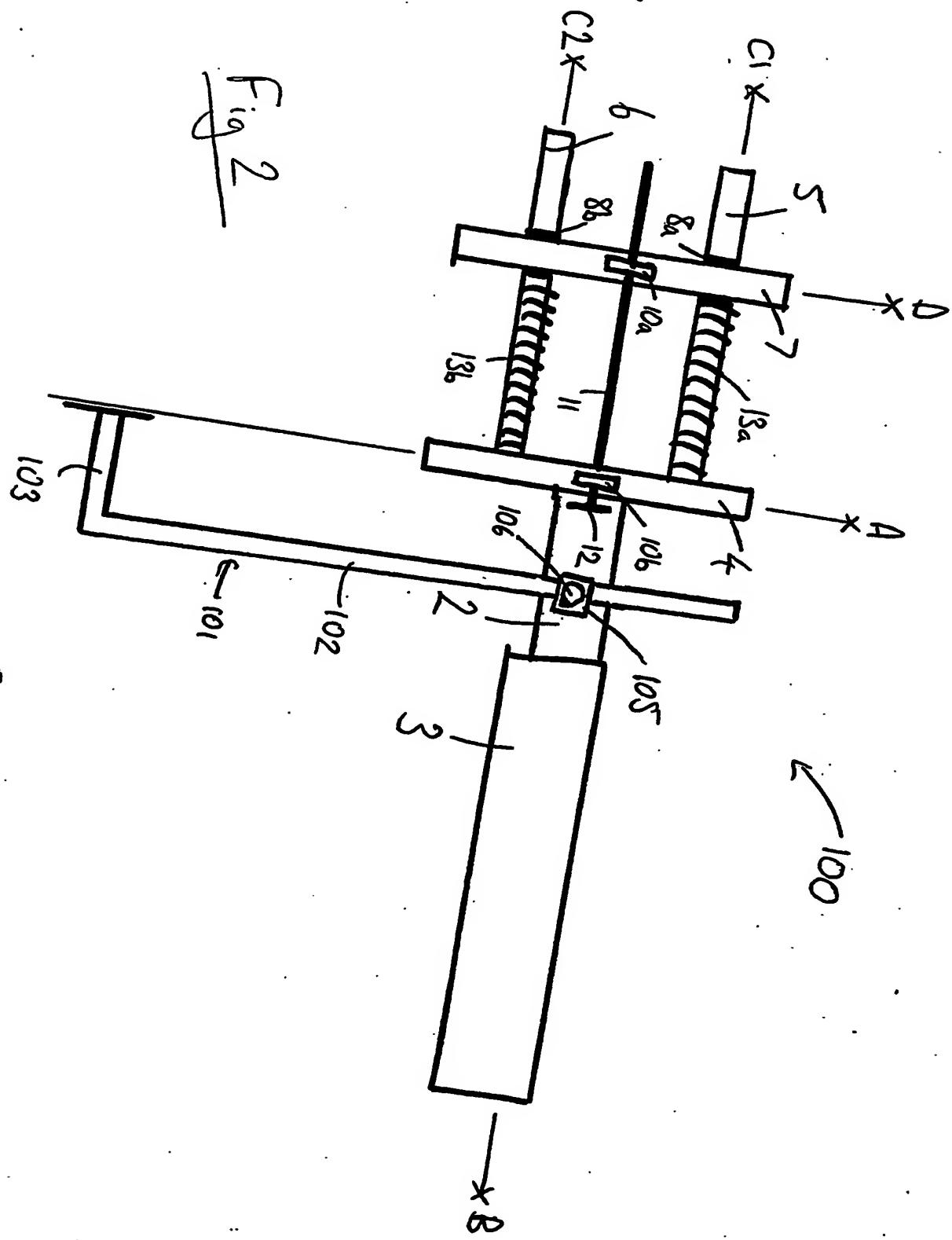
5/15

Fig 1c



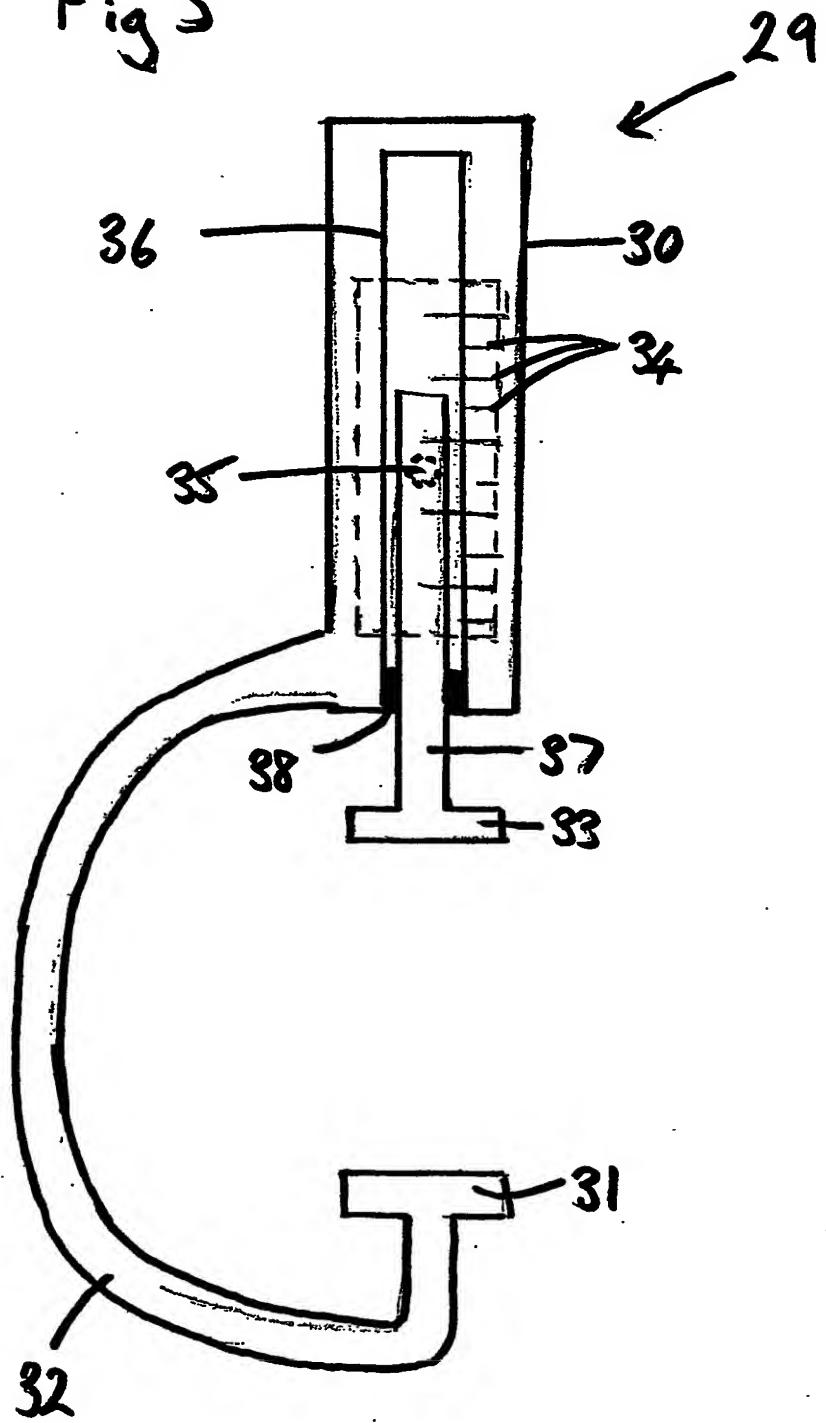
415

Fig 2



515

Fig 3



Document made available under the Patent Cooperation Treaty (PCT)

International application number: PCT/GB04/005168

International filing date: 09 December 2004 (09.12.2004)

Document type: Certified copy of priority document

Document details: Country/Office: GB
Number: 0419983.2
Filing date: 09 September 2004 (09.09.2004)

Date of receipt at the International Bureau: 02 March 2005 (02.03.2005)

Remark: Priority document submitted or transmitted to the International Bureau in compliance with Rule 17.1(a) or (b)



World Intellectual Property Organization (WIPO) - Geneva, Switzerland
Organisation Mondiale de la Propriété Intellectuelle (OMPI) - Genève, Suisse

**This Page is Inserted by IFW Indexing and Scanning
Operations and is not part of the Official Record**

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- BLACK BORDERS**
- IMAGE CUT OFF AT TOP, BOTTOM OR SIDES**
- FADED TEXT OR DRAWING**
- BLURRED OR ILLEGIBLE TEXT OR DRAWING**
- SKEWED/SLANTED IMAGES**
- COLOR OR BLACK AND WHITE PHOTOGRAPHS**
- GRAY SCALE DOCUMENTS**
- LINES OR MARKS ON ORIGINAL DOCUMENT**
- REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY**
- OTHER:** _____

IMAGES ARE BEST AVAILABLE COPY.

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.